In the Claims

Please amend Claims 1, 2, 7, 17, 18, 20, 22, 23 and 29, as follows.

- 1 1. (Currently Amended). An orbital implant which comprises:
- 2 a porous core;
- 3 [[a]] an anterior first coating portion covering a first outer surface section of said core; and
- said first coating portion having a first bioabsorbability rate; and
- a second coating portion, distinct from said first portion, covering a second outer surface
- 6 section of said core; said second coating portion having a second bioabsorbability rate different from
- 7 said first bioabsorbability rate.
- 2. (Currently Amended). The implant of Claim 1, wherein said coating [[is]] portions are deformed
- 2 to intimately contact surface features on said core.
- 1 3. (Previously Presented). The implant of Claim 1, wherein at least one of said coating portions
- 2 comprises a polymer.
- 4. (Previously Presented). The implant of Claim 3, wherein said polymer comprises a material
- 2 selected from the group consisting of polyglycolic acid, polylactic acid, polycaprolactone,
- 3 polydiox-anone, polycyanoacrylate, polyorthoester, poly(gamma-ethyl glutamate), and pseudo-poly
- 4 (amino acid).

- 5. (Previously Presented). The implant of Claim 1, wherein at least one of said coating portions
- 2 comprises a therapeutic agent.
- 6. (Previously Presented). The implant of Claim 5, wherein said therapeutic agent is selected from
- the group consisting of a vascularization agent, and antibiotic agent, an immuno-suppressant, a
- 3 wound-healing promoter, a blood-clot dissolving agent, a blood-clotting agent, a cell-adhesion
- 4 modulating molecule, and any combination thereof.
- 7. (Currently Amended). The implant of Claim 1, wherein said first and second coating portions
- 2 are bonded to one another along a bond.
- 8. (Previously Presented). The implant of Claim 7, wherein said bond is selected from the group
- 2 consisting of: glued bonds, chemical bonds, molecular bonds, magnetic bonds, electrostatic bonds,
- 3 ultrasonic welds, heat welds, press fittings, snap fittings, shrink fittings, friction fittings, and
- 4 mechanically fastened bonds.
- 9. (Previously Presented). The implant of Claim 1, wherein at least one of said coating portions
- 2 comprises a first material having a thickness selected to allow melting penetration using a handheld
- 3 cautery.
- 1 10. (Previously Presented). The implant of Claim 1, which further comprises an indicia identifying
- 2 said first portion.

- 1 11. (Withdrawn). The implant of Claim 10, wherein said indicia comprises lettering.
- 1 12. (Previously Presented). The implant of Claim 10, wherein said indicia comprises a color
- 2 coding.
- 1 13. (Previously Presented). The implant of Claim 1, wherein at least one of said coating portions
- 2 has a passageway therethrough.
- 1 14. (Previously Presented). The implant of Claim 13, wherein said passageway is positioned on a
- 2 posterior location of said implant.
- 1 15. (Previously Presented). The implant of Claim 13, wherein said passageway is sized to allow
- 2 fluid exchange therethrough.
- 1 16. (Previously Presented). The implant of Claim 13, wherein said passageway has an upper rim
- at the surface of said coating portion, and a portion of said core extends into said passageway up to
- a buffer distance from said upper rim.
- 1 17. (Currently Amended). The implant of Claim 1, wherein said first <u>coating</u> portion comprises
- 2 two concentrically adjacent layers wherein a first of said layers comprises a material not present in
- a second of said layers.

- 18. (Currently Amended). The implant of Claim 1, wherein at least one of said coating portions 1 2 comprises means for reducing an adverse immune response by a recipient an immunosuppressant 3 agent. 19. (Previously Presented). The implant of Claim 1, wherein said coating portions have a thickness 1 2 of less than one millimeter. 20. (Currently Amended). An orbital implant which comprises: 1 2 an implant having an outer first surface; a coating at least partially covering said first surface; 3 said coating having a first exposed portion having a first bioabsorbability rate and a separate 4 5 second exposed portion, distinct from said first portion, having a second bioabsorbability rate 6 different from said first bioabsorbability rate.
- 21. (Original). The implant of Claim 20, wherein said coating has an outer second surface which is smoother than said first surface.
- 1 22. (Currently Amended). An orbital implant comprising:
- a substantially spheroid body sized and shaped to be placed in the orbit;
- a coating sized and shaped to intimately contact a section of said body; and
- 4 wherein said coating has a first portion having a first bioabsorbability rate and a separate

- second portion, distinct from said first portion, having a second bioabsorbability rate different from
- 6 said first bioabsorbability rate.
- 1 23. (Currently Amended). The implant of Claim 22, wherein said coating comprises means for
- 2 reducing an adverse immune response by a recipient an immunosuppressant agent.
- 1 24. (Original). The implant of Claim 22, wherein said coating comprises a polymer.
- 1 25. (Previously Presented). The implant of Claim 24, wherein said polymer comprises a material
- 2 selected from the group consisting of polyglycolic acid, polylactic acid, polycaprolactone,
- 3 polydiox-anone, polycyanoacrylate, polyorthoester, poly(gamma-ethyl glutamate), and pseudo-poly
- 4 (amino acid).
- 1 26. (Original). The implant of Claim 22, wherein said coating comprises a therapeutic agent.
- 1 27. (Previously Presented). The implant of Claim 26, wherein said therapeutic agent is selected
- from the group consisting of a vascularization agent, and antibiotic agent, an immuno-suppressant,
- a wound-healing promoter, a blood-clot dissolving agent, a blood-clotting agent, a cell-adhesion
- 4 modulating molecule, and any combination thereof.
- 1 28. (Original). The implant of Claim 22, wherein said coating comprises a surface having
- 2 microtexturing.

1	29. (Currently Amended). A combination of a body and a coating for implantation into the orbit of
2	a mammal;
3	said body comprises an arcuate outer surface;
4	said coating comprises:
. 5	a first external portion being made from a first material having a first
6	bioabsorbability property;
7	said first portion being sized and shaped to intimately contact said outer surface;
8	a second external portion, separate and distinct from said first portion, being made
9	from a second material having a second bioabsorbability property;
10	said second portion being sized and shaped to intimately contact said outer surface;
11	wherein said first bioabsorbability property is different from second bioabsorbability
. 12	property.

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